MANUFACTURER’S DECLARATION OF CONFORMITY
AUSTRALIAN THERAPEUTIC GOODS (MEDICAL DEVICES) REGULATIONS 2002
DECLARATION OF CONFORMITY PROCEDURES

SAP DIR No.: 80021358 Version: B

This is a declaration of conformity made under clause 6.6 of Schedule 3 to the Therapeutic Goods (Medical Devices) Regulations 2002.

Manufacturer’s name: Welch Allyn, Inc.
Business address: 4341 State Street Road
Skaneateles Falls, NY 13153
U.S.A.

Product name: EAR SPECULA for use with Welch Allyn Otoscopes

REF
901001 ACCESSORY, EYE, EAR, NOSE & THROAT

#
22002, 22003, 22004, 22009, 22023, 22025, 22100, 22120, 24302-U,
24303-U, 24304-U, 24305-U, 24320, 24320-B, 24323, 24325, 24327,
24330, 24330-B, 24400-U, 52133, 52134, 52135, 52432-U, 52434-U,
52432-UB, 52434-UB.

Classification: I
GMDN code and term: 35348 – Speculum, ear
Scope of application: All

Each kind of medical device to which the technical documentation applies complies with the applicable provisions of the essential principles and the classification rules before being supplied.

Standards applied: EN ISO 10993-1 2009 + Corr 2010
Biological evaluation of medical devices - Part 1: Evaluation and testing within a risk management process

Authorised Signatory:

Joshua Kim Senior Manager, Regulatory Affairs 2016-08-10
Skaneateles Falls, NY USA Place of Issue

This authorisation is given in the signatory’s capacity as representative of the “Manufacturer” (as recorded above in this declaration)